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# Possibilities and obstacles in recovery of genomes from elusive microbes in complex metagenomes

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## Introduction

Representative genomes provide an entry point for understanding a given microbial ecosystem. The genomes give insights into the metabolic potential and possible roles of the bacteria, as well as being essential when applying other -omics based techniques. For elusive, unculturable bacteria, metagenomics can be a useful tool to extract their genomes directly from their environments. However the approach has limitations when the community complexity is high and microdiversity is present.

In this study, we explored the potentials and obstacles faced when assembling genomes from complex metagenomes using activated sludge as a model system.

## Aims

- To investigate the impact of microdiversity and community complexity on metagenomic assembly, binning and genome extraction.
- To investigate whether short-term enrichment can mitigate the adverse effects of the above

## Methods

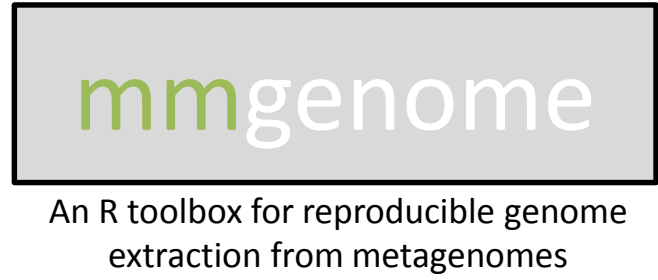
### Short-term Enrichment

- Seed sample**
  - Fullscale EBPR WWTP
- Batch Incubations**
  - Casein, Glucose or Butyrate
  - Aerobic or Anaerobic
  - All incubations at 20 °C
  - 1 sample/d for 7 days
- Screen incubations**
  - V4 16S rRNA amplicons
  - 14 of 50+ samples chosen for metagenome sequencing



### Metagenome Binning

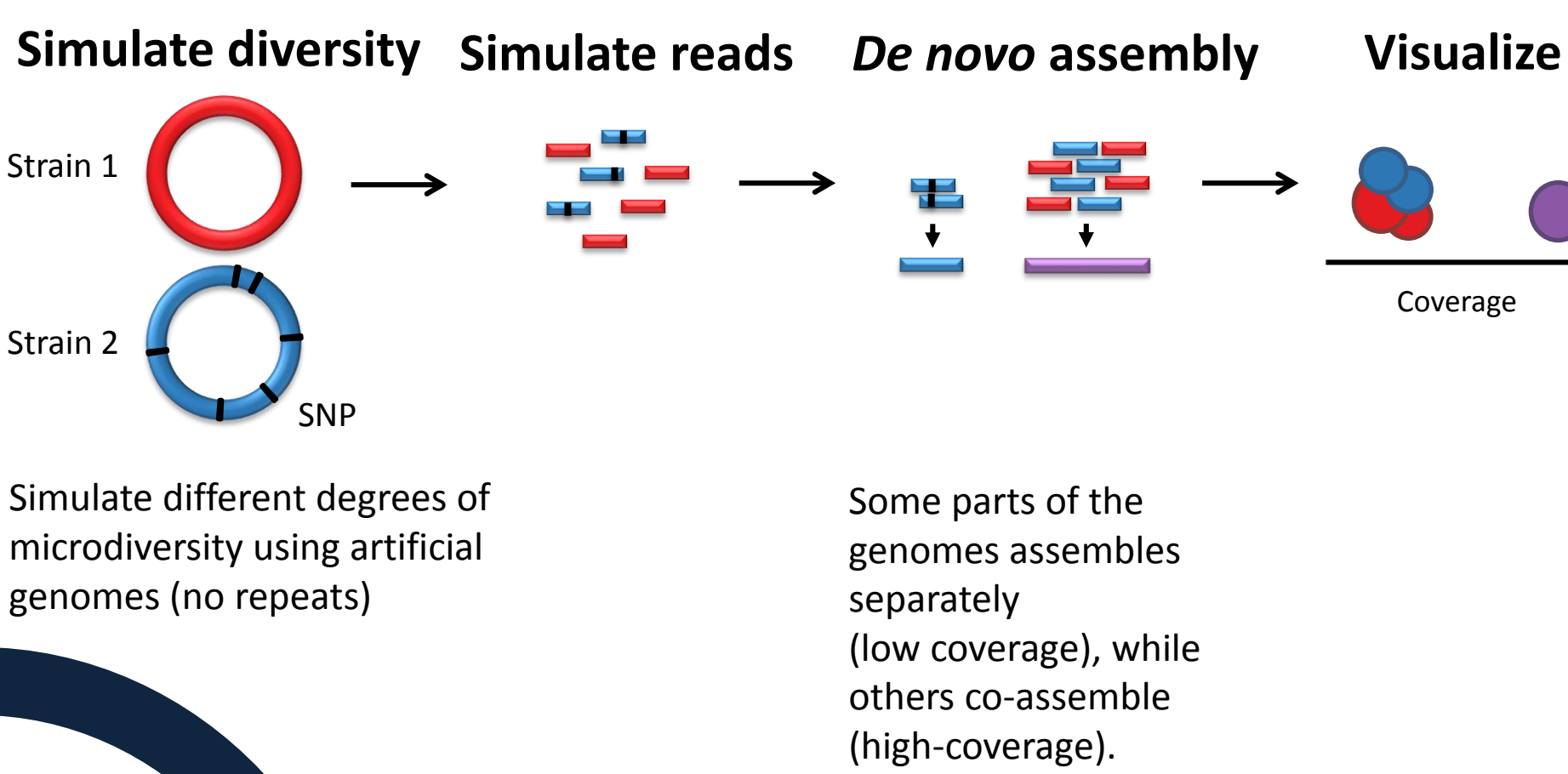
- Sequencing and Assembly**
  - Illumina PE/MP 2x150bp (300 Gbp)
  - De novo assembly (kmer: 64) and read mapping using CLC genomics workbench
- Coverage Binning**
  - Read coverage profiles from 14 samples
  - Taxonomic assignments
  - Essential genes
  - Paired-end read connections
  - Tetranucleotide frequencies



- Re-Assembly and Finishing**
  - Bin specific reads de novo re-assembled in SPAdes
  - Manual finishing using SSPACE, Circo and R-biostrings



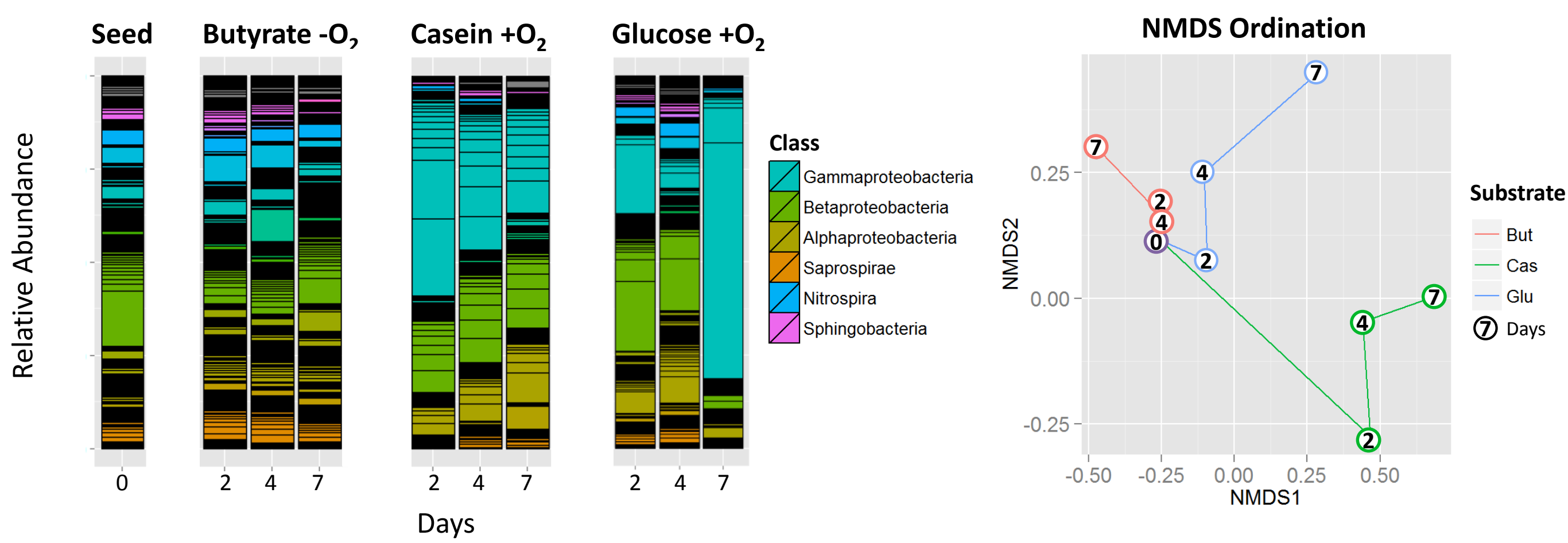
### Microdiversity Simulation



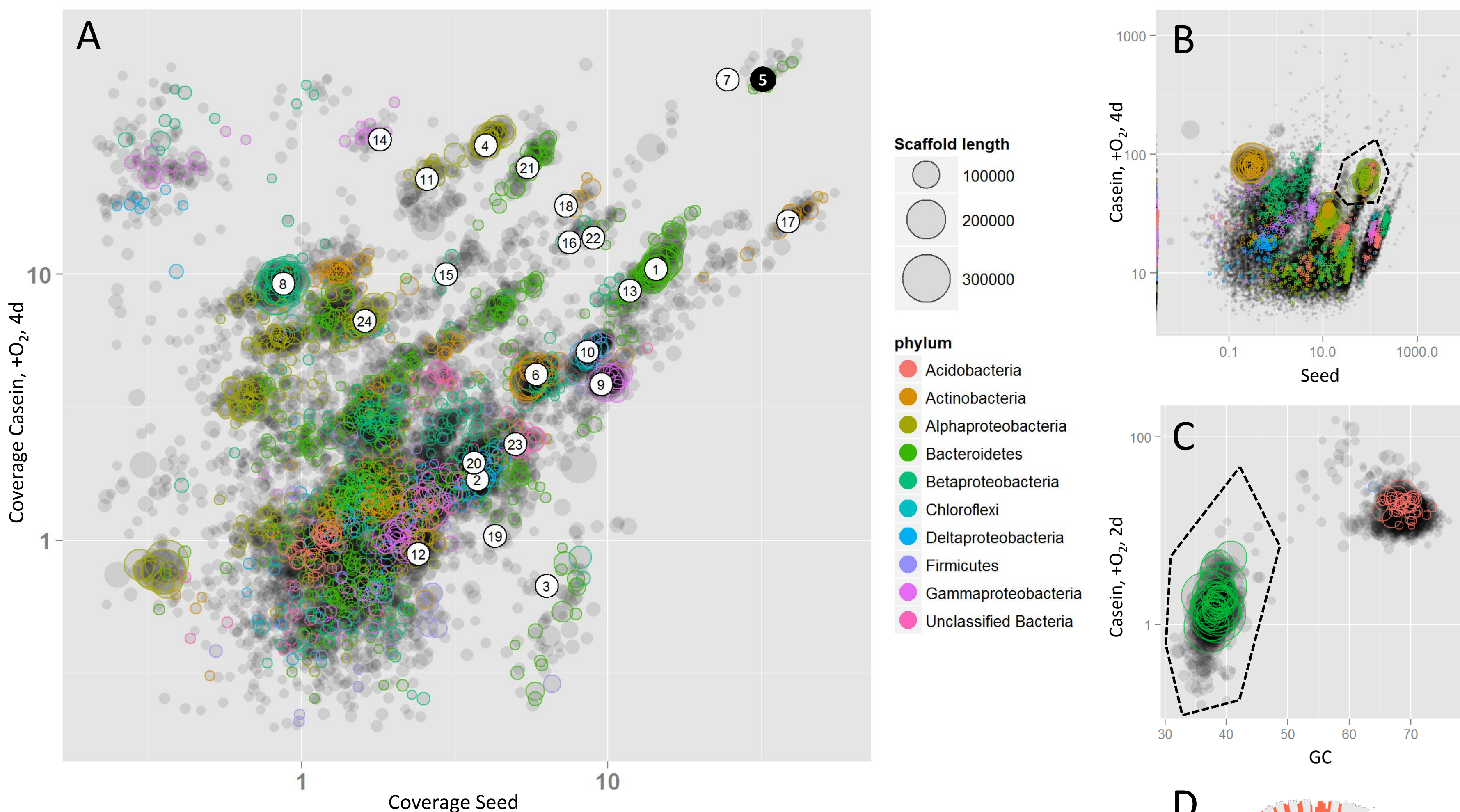
## Conclusions

- Enrichments decreased overall complexity, changed relative abundance and facilitated better coverage binning and allowed for numerous high quality genome bins to be extracted.
- Despite the short-term enrichments microdiversity was still present, which compromised recovery of complete genomes from many species.
- The simulation underlines the devastating impact of micro-diversity on assembly and binning.

## Results



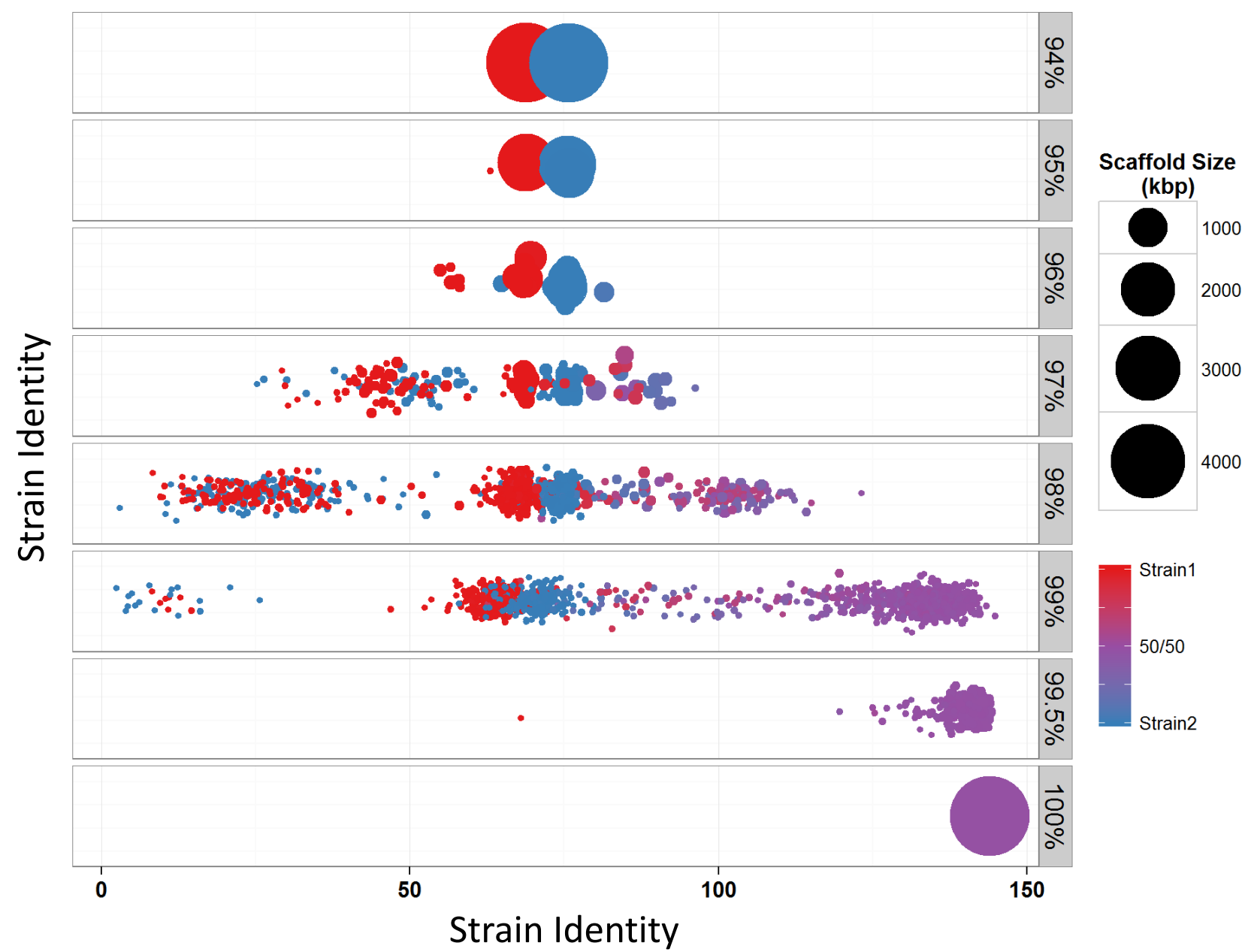
**Screening of short-term enrichments with 16S V4 Amplicon sequencing.** Data shown for 3 of 6 incubations. The 16S screening was used to select samples for further metagenome sequencing.



**Differential coverage binning and Re-assembly** Points represents scaffolds, scaled by their length and colored based on the presence of essential single copy genes. (A) Is a coverage plot of a metagenome assembly of all data. Circled numbers indicate extracted high quality genome bins. In this example bin 5 (A) was targeted for extraction. Bin 5 was found to be enriched in the casein incubation and a sample from this was therefore used for targeted metagenome re-assembly. Bin 5 was binned again from the casein sample metagenome (B & C), and subsequently refined with re-assembly and scaffolding (D). The extracted genome belongs to phylum *Bacteroidetes* and genus *Ferruginibacter*.

	All data		Casein, +O2, 4d		Re-assembly
	Metagenome	Bin 5	Metagenome	Bin 5	Bin 5
Size (Mbp)	2048.3	2.7	125.9	3	3.1
Scaffolds	915 040	702	62422	357	21
Scaffold Max (kbp)	340	34	94	67	1382
Ess. Genes	43522	93/105	2996	104/105	105/105
Species	> 430	1	>29	1	1
Seq. data (Gbp)	300	-	15.5	-	-

**Binning Statistics.** The statistics show the genome extraction progression from the metagenome assemblies, through binning to the final refinement. It is clear that the assembly of the enriched casein sample with less microdiversity provides a better basis for binning, despite far less sequence data being used.



**Simulation of different degrees of microdiversity.** Strain identity refers to the percent bp in the two artificial genomes that are identical.

